Eligible Professional Meaningful Use Menu Set Measures Measure 10 of 10

Stage 1

Date issued: November 7, 2010

Syndromic Surveillance Data Submission	
Objective	Capability to submit electronic syndromic surveillance data to public health agencies and actual submission according to applicable law and practice.
Measure	Performed at least one test of certified EHR technology's capacity to provide electronic syndromic surveillance data to public health agencies and follow-up submission if the test is successful (unless none of the public health agencies to which an EP submits such information has the capacity to receive the information electronically).
Exclusion	An EP who does not collect any reportable syndromic information on their patients during the EHR reporting period or does not submit such information to any public health agency that has the capacity to receive the information electronically.

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Definition of Terms

Public Health Agency -- An entity under the jurisdiction of the U.S. Department of Health and Human Services, tribal organization, State level and/or city/county level administration that serves a public health function.

Attestation Requirements

YES / NO / EXCLUSION

- Eligible professionals (EPs) must attest YES to having performed at least one test of certified EHR technology's capacity to submit electronic syndromic surveillance data to public health agencies and follow up submission if the test was successful (unless none of the public health agencies to which the EP submits such information has the capacity to receive the information electronically) to meet this measure.
- EXCLUSION: If an EP does not collect any reportable syndromic information on their patients during the EHR reporting period or if no public health agency that has the capacity to receive the information electronically, then the EP is excluded from this requirement. EPs must select NO next to the appropriate exclusion, then click the APPLY button in order to attest to the exclusion.

Additional Information

- The test to meet the measure of this objective must involve the actual submission of electronic syndromic surveillance data to public health agencies, if one exists that will accept the information. Simulated transfers of information are not acceptable to satisfy this objective.
- The transmission of electronic syndromic surveillance data is not required for the purposes of a test. The use of test information about a fictional patient that would be identical in form to what would be sent about an actual patient would satisfy this objective.
- An unsuccessful test to submit electronic syndromic surveillance data to public health agencies will be considered valid and would satisfy this objective.
- If the test is successful, then the EP should institute regular reporting with the entity with whom the successful test was conducted, in accordance with applicable law and practice. There is not a measurement associated with this reporting.
- EPs must test their ability to submit electronic syndromic surveillance data to public health agencies at least once prior to the end of the EHR reporting period. Testing may also occur prior to the beginning of the EHR reporting period. Each payment year requires it own unique test.
- If multiple EPs are using the same certified EHR technology in a shared physical setting, testing would only have to occur once for a given certified EHR technology.
- The transmission of syndromic surveillance information must use the standards at 45 CFR 170.302(l).

Related FAQs

- **#10764** - To meet the public health meaningful use objectives (submitting information to an immunization registry, reporting lab results to a public health agency, or reporting syndromic surveillance information), does a provider have to send information directly from their certified EHR technology to the appropriate receiving entity or can they use an intermediary such as a HIE or another third-party software vendor?
- **#10714** - If my certified EHR technology only includes the capability to submit information to an immunization registry using the HL7 2.3.1 standard but the immunization registry only accepts information formatted in the HL7 2.5.1 or some other standard, will I qualify for an exclusion because the immunization registry does not have the capacity to receive the information electronically? What if the immunization registry has a waiting list or is unable to test for other reasons but can accept information formatted in HL7 2.3.1, is that still a valid exclusion?
- **#10532** - Will the requirement that EPs and eligible hospitals choose at least one public health objective among the meaningful use measures still apply to those States that ask CMS for approval to change the definition of meaningful use?
- **#10151** - If an EP is unable to meet the measure of a meaningful use objective because it is outside of the scope of his or her practice, will the EP be excluded from meeting the measure of that objective?
- **#10162** - How should EPs select menu objectives?